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* * * * * * * * * *   Welcome to STN International   * * * * * * * * * *

 NEWS  1           Web Page URLs for STN Seminar Schedule - N. America
 NEWS  2           "Ask CAS" for self-help around the clock
 NEWS  3  SEP 09   CA/CAplus records now contain indexing from 1907 to the
                   present
 NEWS  4  AUG 05   New pricing for EUROPATFULL and PCTFULL effective
                   August 1, 2003
 NEWS  5  AUG 13   Field Availability (/FA) field enhanced in BEILSTEIN
 NEWS  6  AUG 18   Data available for download as a PDF in RDISCLOSURE
 NEWS  7  AUG 18   Simultaneous left and right truncation added to PASCAL
 NEWS  8  AUG 18   FROSTI and KOSMET enhanced with Simultaneous Left and Righ
                   Truncation
 NEWS  9  AUG 18   Simultaneous left and right truncation added to ANABSTR
 NEWS 10  SEP 22   DIPPR file reloaded
 NEWS 11  SEP 25   INPADOC: Legal Status data to be reloaded
 NEWS 12  SEP 29   DISSABS now available on STN
 NEWS 13  OCT 10   PCTFULL: Two new display fields added
 NEWS 14  OCT 21   BIOSIS file reloaded and enhanced
 NEWS 15  OCT 28   BIOSIS file segment of TOXCENTER reloaded and enhanced
 NEWS 16  NOV 24   MSDS-CCOHS file reloaded

 NEWS EXPRESS  NOVEMBER 14 CURRENT WINDOWS VERSION IS V6.01c, CURRENT
               MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP),
               AND CURRENT DISCOVER FILE IS DATED 23 SEPTEMBER 2003
 NEWS HOURS    STN Operating Hours Plus Help Desk Availability
 NEWS INTER    General Internet Information
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 NEWS PHONE    Direct Dial and Telecommunication Network Access to STN
 NEWS WWW      CAS World Wide Web Site (general information)

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FILE 'HOME' ENTERED AT 14:28:10 ON 01 DEC 2003

=> file registry
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                       ENTRY    SESSION
FULL ESTIMATED COST                                     0.21       0.21

FILE 'REGISTRY' ENTERED AT 14:28:19 ON 01 DEC 2003
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STRUCTURE FILE UPDATES:   30 NOV 2003  HIGHEST RN 622330-21-6
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DICTIONARY FILE UPDATES: 30 NOV 2003 HIGHEST RN 622330-21-6

TSCA INFORMATION NOW CURRENT THROUGH JULY 14, 2003

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details:
http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf

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L1 STRUCTURE UPLOADED

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L1 HAS NO ANSWERS
'STR ' IS NOT A VALID STRUCTURE FORMAT KEYWORD
Structure Formats
SIA ----- Structure Image, Attributes, and map table if it contains data. (Default)
SIM ----- Structure IMage.
SAT ----- Structure ATtributes and map table if it contains data.
SCT ----- Structure Connection Table and map table if it contains data.
SDA ----- All Structure DAta (image, attributes, connection table and map table if it contains data).
NOS ----- NO Structure data.
ENTER STRUCTURE FORMAT (SIA), SCT, SDA, SIM, SAT, NOS:end

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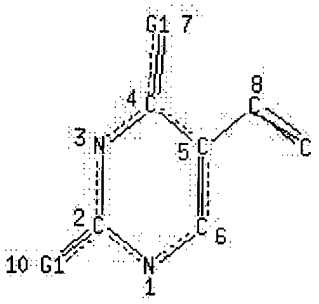

Page 1-A

9

Page 1-B
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NSPEC IS R AT 2
NSPEC IS R AT 3
NSPEC IS R AT 4

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DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
RING(S) ARE ISOLATED OR EMBEDDED
NUMBER OF NODES IS  12

STEREO ATTRIBUTES: NONE


=> s l1 full
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FULL SCREEN SEARCH COMPLETED -   2151 TO ITERATE

100.0% PROCESSED    2151 ITERATIONS                          1211 ANSWERS
SEARCH TIME: 00.00.01

L2         1211 SEA SSS FUL L1

=> file caplus
COST IN U.S. DOLLARS                          SINCE FILE      TOTAL
                                                 ENTRY    SESSION
FULL ESTIMATED COST                             148.55     148.76

FILE 'CAPLUS' ENTERED AT 14:29:26 ON 01 DEC 2003
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FILE COVERS 1907 - 1 Dec 2003  VOL 139 ISS 23
FILE LAST UPDATED: 30 Nov 2003  (20031130/ED)

  This file contains CAS Registry Numbers for easy and accurate
  substance identification.


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L4         121 L3 AND PY<1992

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L5          45 L4 AND (DNA OR NUCLEIC)

=> d 1-45 ti

L5   ANSWER 1 OF 45  CAPLUS  COPYRIGHT 2003 ACS on STN
TI   Relationships between nucleotide incorporation rates and molecular
     parameters obtained by molecular modeling and chromatography

L5   ANSWER 2 OF 45  CAPLUS  COPYRIGHT 2003 ACS on STN
TI   Preparation of fluorescence-labeled DNA by polymerase chain reaction (PCR)

L5   ANSWER 3 OF 45  CAPLUS  COPYRIGHT 2003 ACS on STN
TI   Antimalarial compositions containing pyrimidine analog inhibitors of
     nucleic acid biosynthesis

L5   ANSWER 4 OF 45  CAPLUS  COPYRIGHT 2003 ACS on STN
TI   Spectrally resolvable rhodamine dyes for nucleic acid sequence
     determination

L5   ANSWER 5 OF 45  CAPLUS  COPYRIGHT 2003 ACS on STN
TI   Nucleic acid related compounds. 65. New syntheses of
     1-(β-D-arabinofuranosyl)-5(E)-(2-iodovinyl)uracil (IVAraU) from
     vinylsilane precursors. Radioiodine uptake as a marker for thymidine
     kinase herpes viral infections

L5   ANSWER 6 OF 45  CAPLUS  COPYRIGHT 2003 ACS on STN
TI   Preparation of crosslinking oligonucleotides as nucleic acid
     hybridization probes

L5   ANSWER 7 OF 45  CAPLUS  COPYRIGHT 2003 ACS on STN
TI   Palladium-catalyzed approach to 5-substituted uracil and uridine
     derivatives

L5   ANSWER 8 OF 45  CAPLUS  COPYRIGHT 2003 ACS on STN
TI   Analysis of mutations in the thymidine kinase genes of drug-resistant
     varicella-zoster virus populations using the polymerase chain reaction

L5   ANSWER 9 OF 45  CAPLUS  COPYRIGHT 2003 ACS on STN
TI   Nucleic acid related compounds. 61. Conversion of vinylsilanes to
     vinyl halides with xenon difluoride and metal halides. A versatile new
     route to 5-(2-halovinyl)pyrimidine nucleosides

L5   ANSWER 10 OF 45  CAPLUS  COPYRIGHT 2003 ACS on STN
TI   Nucleic acid related compounds. 59. Solvent, not palladium oxidation
     state, is the primary determinant for successful coupling of terminal
     alkynes with iodonucleosides

L5   ANSWER 11 OF 45  CAPLUS  COPYRIGHT 2003 ACS on STN
TI   DNA dynamics from a spin probe: dependence of probe motion on tether
     length

L5   ANSWER 12 OF 45  CAPLUS  COPYRIGHT 2003 ACS on STN
TI   Application of chromatographic retention data in an investigation of a
     quantitative structure-nucleotide incorporation rate relationship

L5   ANSWER 13 OF 45  CAPLUS  COPYRIGHT 2003 ACS on STN
TI   Palladium-catalyzed alkylations in aqueous media

L5   ANSWER 14 OF 45  CAPLUS  COPYRIGHT 2003 ACS on STN
TI   Automated DNA sequence analysis and DNA fingerprinting
```

L5 ANSWER 15 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Mapping restriction sites on **DNA** with fluorescent labels and interrupted-palindrome restriction enzymes

L5 ANSWER 16 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Correlation of nucleotide incorporation rate and HPLC retention parameters of substituted nucleosides

L5 ANSWER 17 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Sequence- and structure-dependent **DNA** base dynamics: synthesis, structure, and dynamics of site, and sequence specifically spin-labeled **DNA**

L5 ANSWER 18 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Palladium-catalyzed synthesis of alkynylamino nucleosides. A universal linker for **nucleic** acids

L5 ANSWER 19 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Effect of manganese ions on the incorporation of dideoxynucleotides by bacteriophage T7 **DNA** polymerase and Escherichia coli **DNA** polymerase I

L5 ANSWER 20 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Preparation of 5-(3-aminoprop-1-ynyl)deoxyuridine derivatives and their use for synthesis of **DNA** and RNA labeled with nonradioactive markers.

L5 ANSWER 21 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI **DNA** structural data from a dynamics probe. The dynamic signatures of single stranded, hairpin-looped, and duplex forms of **DNA** are distinguishable

L5 ANSWER 22 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Design and synthesis of fluorescently labeled chain terminators for automated sequencing of **DNA**

L5 ANSWER 23 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Preparation of (aminoalkynyl)nucleotides as intermediates for fluorescent chain terminators for **DNA** sequencing

L5 ANSWER 24 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI **DNA** sequencing by fluorescence analysis

L5 ANSWER 25 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Preparation of base-modified nucleosides suitable for non-radioactive label attachment and their incorporation into synthetic oligodeoxyribonucleotides

L5 ANSWER 26 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Potent activity of 5-fluoro-2'-deoxyuridine and related compounds against thymidine kinase-deficient (TK-) herpes simplex virus: targeted at thymidylate synthase

L5 ANSWER 27 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI A rigid and nonperturbing probe for duplex **DNA** motion

L5 ANSWER 28 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Synthesis and application of derivatizable oligonucleotides

L5 ANSWER 29 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI A system for rapid **DNA** sequencing with fluorescent chain-terminating dideoxynucleotides

L5 ANSWER 30 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Substrate specificity of **DNA** polymerases. II. 5-(1-Alkynyl)-dUTPs as substrates of the Klenow **DNA** polymerase enzyme

L5 ANSWER 31 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI The synthesis and biological properties of some 5-substituted-2'-deoxyuridines

L5 ANSWER 32 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Rapid determination of thymidylate synthase activity and its inhibition in intact L1210 leukemia cells in vitro

L5 ANSWER 33 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Incorporation of 5-substituted pyrimidine nucleoside analogs into **DNA** of a thymidylate synthetase-deficient murine FM3A carcinoma cell line

L5 ANSWER 34 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Strategies for the measurement of the inhibitory effects of thymidine analogs on the activity of thymidylate synthase in intact murine leukemia L1210 cells

L5 ANSWER 35 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Thymidylate synthetase-deficient mouse FM3A mammary carcinoma cell line as a tool for studying the thymidine salvage pathway and the incorporation of thymidine analogs into host cell **DNA**

L5 ANSWER 36 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Induction of sister-chromatid exchange by 5-substituted 2'-deoxyuridines

L5 ANSWER 37 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI **Nucleic** acid related compounds. 39. Efficient conversion of 5-iodo to 5-alkynyl and derived 5-substituted uracil bases and nucleosides

L5 ANSWER 38 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI **Nucleic** acid related compounds. 40. Synthesis and biological activities of 5-alkynyluracil nucleosides

L5 ANSWER 39 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI **Nucleic** acid related compounds. 31. Smooth and efficient palladium-copper catalyzed coupling of terminal alkynes with 5-iodouracil nucleosides

L5 ANSWER 40 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Thymidylate synthetase as target enzyme for the inhibitory activity of 5-substituted 2'-deoxyuridines on mouse leukemia L1210 cell growth

L5 ANSWER 41 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Effects of E-5-(2-bromovinyl)-2'-deoxyuridine and other selective antiherpes compounds on the induction of retrovirus particles in mouse BALB/3T3 cells

L5 ANSWER 42 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Comparative study of the potency and selectivity of antiherpes compounds

L5 ANSWER 43 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Antiviral **nucleic** acid derivatives. III. Crystal structure of 5-ethynyl-2'-deoxyuridine

L5 ANSWER 44 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI Incorporation of 5-substituted uracil derivatives into **nucleic** acids. Part IV. The synthesis of 5-ethynyluracil

L5 ANSWER 45 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
TI The preparation and properties of some 5-substituted uracil derivatives

=> d 23-25, 2, 6, 20 bib ab hitstr

L5 ANSWER 23 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
Full Text
AN 1988:493540 CAPLUS
DN 109:93540
TI Preparation of (aminoalkynyl)nucleotides as intermediates for fluorescent chain terminators for **DNA** sequencing
IN Hobbs, Frank Worden, Jr.; Cocuzza, Anthony Joseph
PA du Pont de Nemours, E. I., and Co., USA
SO Eur. Pat. Appl., 40 pp.
CODEN: EPXXDW
DT Patent
LA English
FAN.CNT 2

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PI	EP 251786	A2	19880107	EP 1987-305844	19870701 <--
	EP 251786	A3	19891206		
	EP 251786	B1	19941130		
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	US 5047519	A	19910910	US 1987-57565	19870612 <--
	CA 1340022	A1	19980901	CA 1987-540946	19870630
	DK 8703375	A	19880103	DK 1987-3375	19870701 <--
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	DK 9300819	A	19930707	DK 1993-819	19930707
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	US 5625081	A	19970429	US 1994-181284	19940113
	US 5558991	A	19960924	US 1994-192915	19940207
	US 5608063	A	19970304	US 1995-412409	19950328

PRAI US 1986-881372 19860702
US 1987-57565 19870612
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JP 1996-221531 19870702
US 1991-780346 19911022
US 1991-780347 19911022
US 1992-981148 19921124
US 1993-981026 19930217
US 1994-181358 19940113

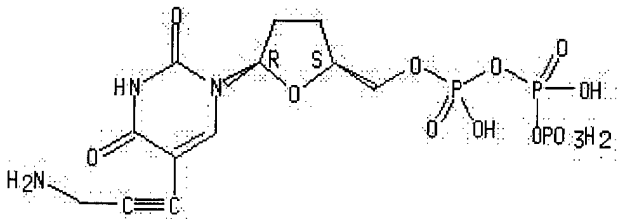
OS MARPAT 109:93540
AB The title compds. [I-IV; R = H, NH2; R1 = R2R3NZC≡C; R2, R3 = H, C1-4 alkyl, protecting group; R4 = sugar moiety Q, Q1, ether moiety Q2; R5 = H, (HO)2P(O), H3P2O6, H4P3O9; when R7 = R8 = H, then R6 = H, OH, F, NH2, N3; when R7 = H, R8 = OH, then R6 = H, OH; when R7 = OH, R8 = H, then R6 = OH; Z = diradical moiety of 1-20 atoms] and their salts were prepd. for coupling with fluorescent dyes to prep. fluorescent chain terminators for **DNA** sequencing. 6-Methoxy-2-(methylthio)-9-(2,3-dideoxy-5-O-trityl-β-D-ribofuranosyl)-7-deazapurine, prepd. in 5 steps from 6-methoxy-2-(methylthio)-7-deazapurine, was iodinated by treatment with

N-iodosuccinimide and the 7-iodo deriv. was de-O-methylated, oxidized to the sulfoxide and ammonolyzed, and detritylated to give azaguanosine IV (R = NH2, R1 = iodo, R4 = Q, R5-R8 = H). This was coupled with HC≡CCH2NHCOCF3 in the presence of (Ph3P)4Pd/CuI catalyst and the product converted to the triphosphate and deprotected to give IV (R = NH2, R1 = H2NCH2C≡C, R4 = Q, R5 = H4P3O9, R6-R8 = H). This was condensed with a xanthene deriv. (prepn. given) to give fluorescent chain terminator V.

IT **114748-59-3P 114748-60-6P 115899-40-6P**
115899-42-8P 115899-44-0P 115899-45-1P
115899-46-2P
RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of, as fluorescent chain terminator intermediate, for **DNA** sequencing)

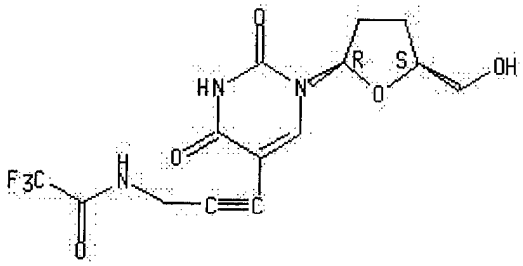
RN 114748-59-3 CAPLUS
CN Uridine 5'-(tetrahydrogen triphosphate), 5-(3-amino-1-propynyl)-2',3'-dideoxy- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



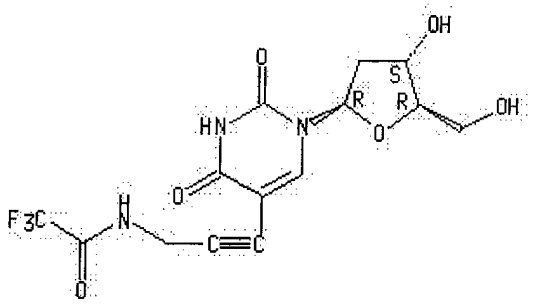
RN 114748-60-6 CAPLUS
CN Uridine, 2',3'-dideoxy-5-[3-[(trifluoroacetyl)amino]-1-propynyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 115899-40-6 CAPLUS
CN Uridine, 2'-deoxy-5-[3-[(trifluoroacetyl)amino]-1-propynyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 115899-42-8 CAPLUS
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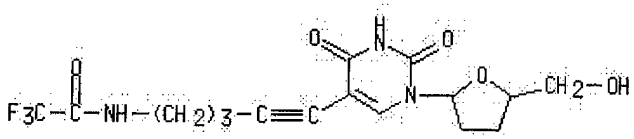
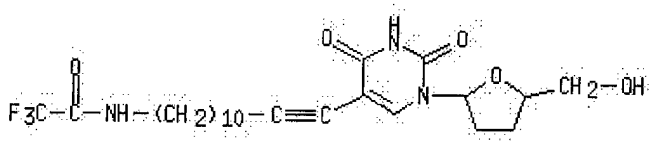
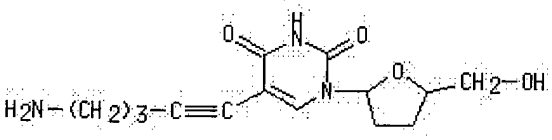


RN 115899-44-0 CAPLUS
CN Uridine, 2',3'-dideoxy-5-[12-[(trifluoroacetyl)amino]-1-dodecynyl]- (9CI) (CA INDEX NAME)

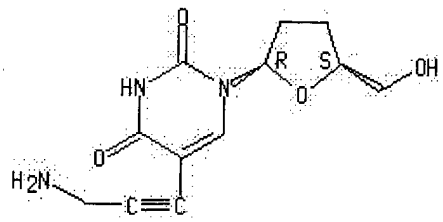


RN 115899-45-1 CAPLUS
CN Uridine, 5-(5-amino-1-pentynyl)-2',3'-dideoxy- (9CI) (CA INDEX NAME)



RN 115899-46-2 CAPLUS
CN Uridine, 5-(3-amino-1-propynyl)-2',3'-dideoxy- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L5 ANSWER 24 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
Full Text
AN 1988:401856 CAPLUS
DN 109:1856
TI **DNA** sequencing by fluorescence analysis
IN Prober, James Merrill; Dam, Rudy Johan; Robertson, Charles William, Jr.; Hobbs, Frank Worden, Jr.; Trainor, George Leonard
PA du Pont de Nemours, E. I., and Co., USA
SO Eur. Pat. Appl., 66 pp.
CODEN: EPXXDW
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 252683	A2	19880113	EP 1987-305848	19870701 <--
	EP 252683	A3	19891123		
	EP 252683	B1	19950118		
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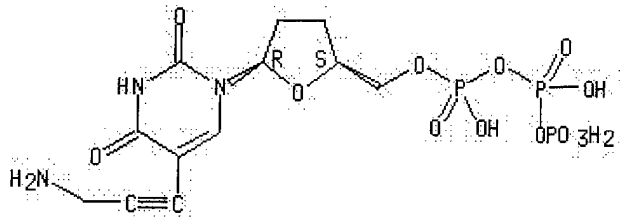
	US 1993-981026		19930217		
	US 1994-181358		19940113		

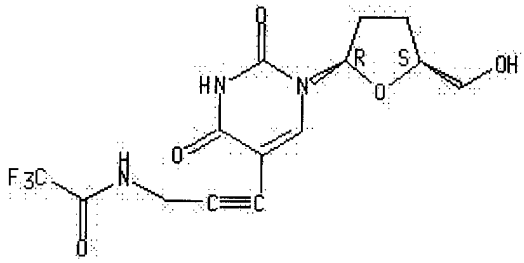
OS MARPAT 109:1856
AB A fluorescence-based system for **DNA** sequence anal. is described. A set of fluorescence-labeled **DNA** chain terminators was generated. Each of the 4 chain terminators corresponding to the 4 nucleotide basis in **DNA** carries a different fluorescent compd. **DNA** fragments to be sequenced can be labeled with these fluorescent compds. in a single vessel. The labeled **DNA** fragments of varying lengths are then sepd. by electrophoresis. A photometric detection system is used to identify the labeled bases and det. the **DNA** sequence.
IT **114748-59-3P 114748-60-6P 114748-72-0P**
RL: PREP (Preparation)
(prepn. of, detn. of **DNA** sequences by fluorescence anal. in relation to)
RN 114748-59-3 CAPLUS
CN Uridine 5'-(tetrahydrogen triphosphate), 5-(3-amino-1-propynyl)-2',3'-dideoxy- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



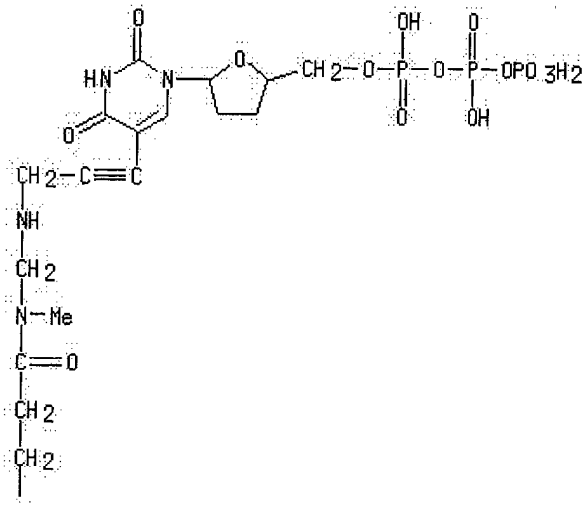
RN 114748-60-6 CAPLUS
CN Uridine, 2',3'-dideoxy-5-[3-[(trifluoroacetyl)amino]-1-propynyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

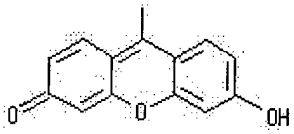


RN 114748-72-0 CAPLUS
CN Uridine 5'-(tetrahydrogen triphosphate), 2',3'-dideoxy-5-[3-[[[[3-(6-hydroxy-3-oxo-3H-xanthen-9-yl)-1-oxopropyl]methylamino]methyl]amino]-1-propynyl]- (9CI) (CA INDEX NAME)

PAGE 1-A

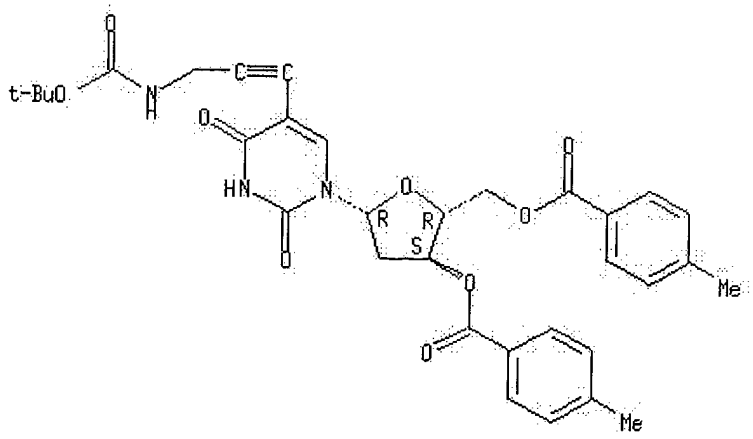


PAGE 2-A



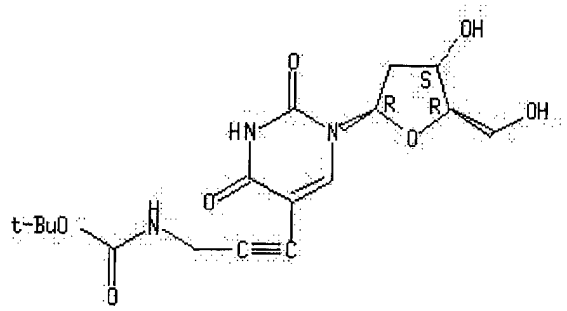
L5 ANSWER 25 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
Full Text
AN 1988:187182 CAPLUS
DN 108:187182
TI Preparation of base-modified nucleosides suitable for non-radioactive label attachment and their incorporation into synthetic oligodeoxyribonucleotides
AU Haralambidis, Jim; Chai, Miao; Tregear, Geoffrey W.
CS Howard Florey Inst. Exp. Physiol. Med., Univ. Melbourne, Parkville, 3052, Australia
SO Nucleic Acids Research (**1987**), 15(12), 4857-76
CODEN: NARHAD; ISSN: 0305-1048
DT Journal
LA English
OS CASREACT 108:187182
AB A very mild and efficient procedure has been developed for the prepn. of C-5 substituted deoxyuridines. The substituent carries a masked primary aliph. amino group. These compds. are readily converted into their phosphoramidites and can be used to prep. oligonucleotides carrying one or more aliph. amino groups. Fluorescein isothiocyanate coupled to these compds. gives oligonucleotide probes carrying multiple fluorescein labels. These compds. have a free 5'-hydroxy group enabling additional 5'- end radioactive labeling for evaluation of their hybridization characteristics. Oligonucleotides carrying a long (11 atom) linker arm to the fluorescein hybridize more efficiently to mRNA than those carrying a short (4 atom) arm. The long linker arm derivs. are comparable to underivatized oligonucleotides in hybridizing to mRNA.
IT **114079-29-7P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. and detoluoylation of)
RN 114079-29-7 CAPLUS
CN Carbamic acid, [3-[1-[2-deoxy-3,5-bis-O-(4-methylbenzoyl)-β-D-erythro-pentofuranosyl]-1,2,3,4-tetrahydro-2,4-dioxo-5-pyrimidinyl]-2-propynyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.



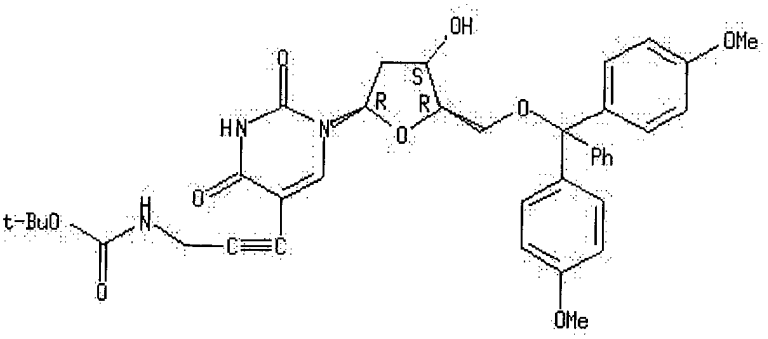
IT **114079-30-0P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. and dimethoxytritylation of)
RN 114079-30-0 CAPLUS
CN Carbamic acid, [3-[1-(2-deoxy-β-D-erythro-pentofuranosyl)-1,2,3,4-tetrahydro-2,4-dioxo-5-pyrimidinyl]-2-propynyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.



IT **114079-31-1P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. and reaction of, with bis(diisopropylamino)methoxyphosphine)
RN 114079-31-1 CAPLUS
CN Carbamic acid, [3-[1-[5-O-[bis(4-methoxyphenyl)phenylmethyl]-2-deoxy-β-D-erythro-pentofuranosyl]-1,2,3,4-tetrahydro-2,4-dioxo-5-pyrimidinyl]-2-propynyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

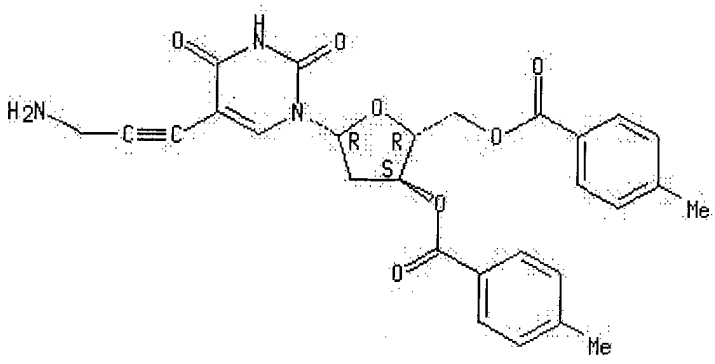


IT **114079-33-3P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. and reaction of, with nitrophenyl aminohexanoate deriv.)
RN 114079-33-3 CAPLUS
CN Uridine, 5-(3-amino-1-propynyl)-2'-deoxy-, 3',5'-bis(4-methylbenzoate), mono(trifluoroacetate) (salt) (9CI) (CA INDEX NAME)

CM 1

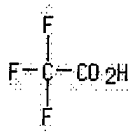
CRN 114079-32-2
CMF C28 H27 N3 O7

Absolute stereochemistry.



CM 2

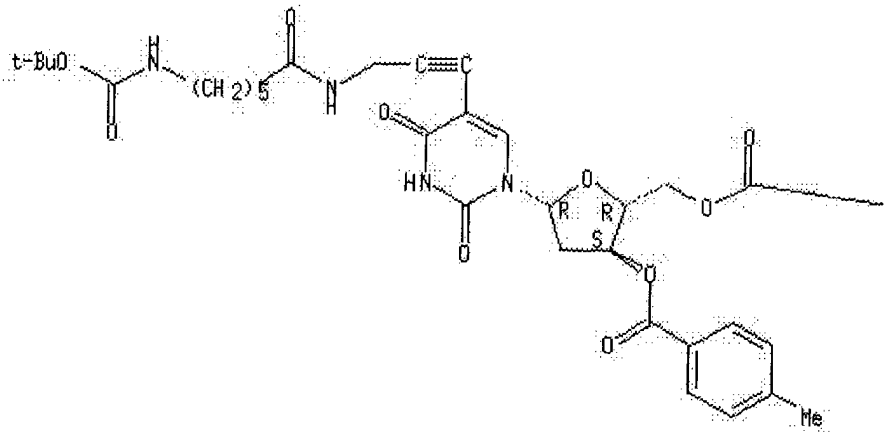
CRN 76-05-1
CMF C2 H F3 O2



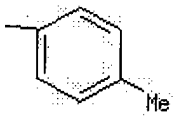
IT **114079-34-4P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. and sequential detoluoylation and dimethoxytritylation of)
RN 114079-34-4 CAPLUS
CN Carbamic acid, [6-[[3-[1-[2-deoxy-3,5-bis-O-(4-methylbenzoyl)-β-D-erythro-pentofuranosyl]-1,2,3,4-tetrahydro-2,4-dioxo-5-pyrimidinyl]-2-propynyl]amino]-6-oxohexyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A



PAGE 1-B



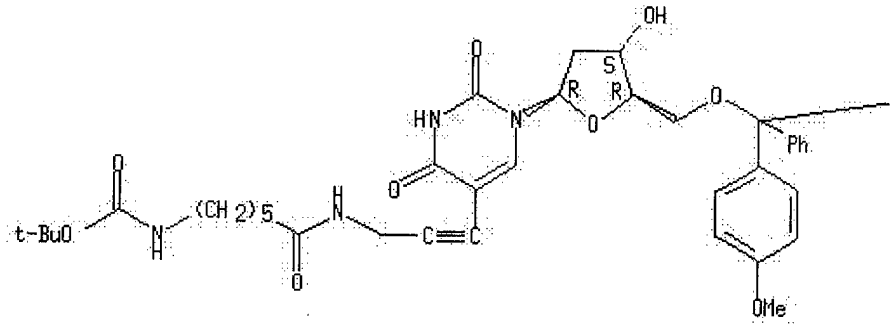
IT **114079-35-5P 114079-36-6P 114103-42-3P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of, intermediate in synthesis of fluorescent-labeled oligodeoxyribonucleotides)

RN 114079-35-5 CAPLUS

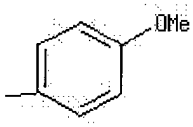
CN Carbamic acid, [6-[[3-[1-[5-O-[bis(4-methoxyphenyl)phenylmethyl]-2-deoxy-β-D-erythro-pentofuranosyl]-1,2,3,4-tetrahydro-2,4-dioxo-5-pyrimidinyl]-2-propynyl]amino]-6-oxohexyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A



PAGE 1-B

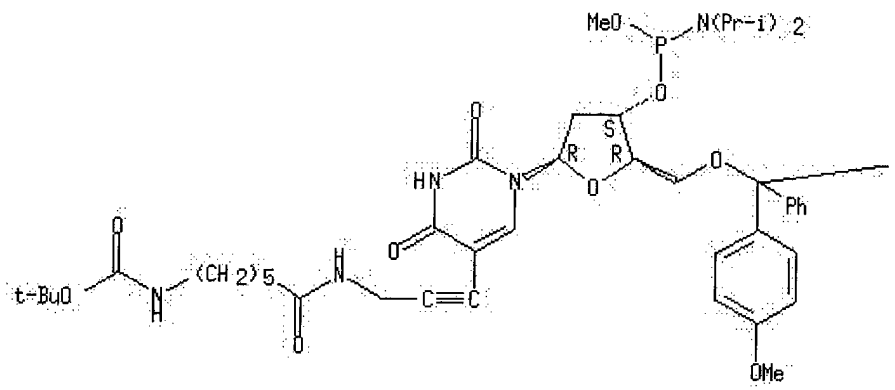


RN 114079-36-6 CAPLUS

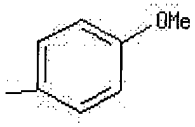
CN Carbamic acid, [6-[[3-[1-[5-O-[bis(4-methoxyphenyl)phenylmethyl]-3-O-[[bis(1-methylethyl)amino]methoxyphosphino]-2-deoxy-β-D-erythro-pentofuranosyl]-1,2,3,4-tetrahydro-2,4-dioxo-5-pyrimidinyl]-2-propynyl]amino]-6-oxohexyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

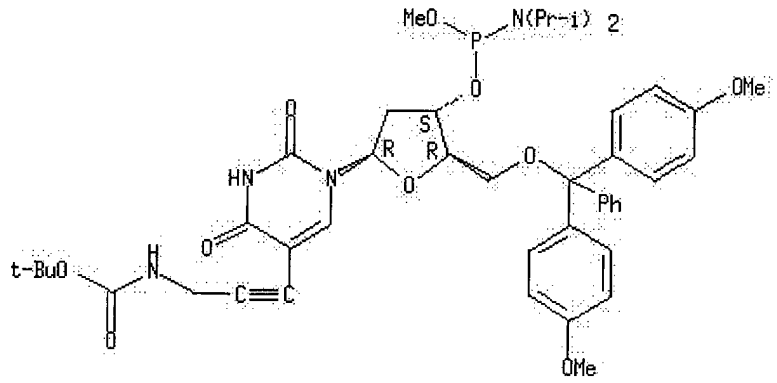


PAGE 1-B



RN 114103-42-3 CAPLUS
CN Carbamic acid, [3-[1-[5-O-[bis(4-methoxyphenyl)phenylmethyl]-3-O-[[bis(1-methylethyl)amino]methoxyphosphino]-2-deoxy-β-D-erythro-pentofuranosyl]-1,2,3,4-tetrahydro-2,4-dioxo-5-pyrimidinyl]-2-propynyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L5 ANSWER 2 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
Full Text
AN 1992:16706 CAPLUS
DN 116:16706
TI Preparation of fluorescence-labeled **DNA** by polymerase chain reaction (PCR)
IN Manabe, Nobuhisa; Uchimura, Yuka; Miyazaki, Keiko; Kato, Ikunoshin
PA Takara Shuzo Co., Ltd., Japan
SO Jpn. Kokai Tokkyo Koho, 6 pp.

CODEN: JKXXAF
DT Patent
LA Japanese
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 03210197	A2	19910913	JP 1990-3437	19900112 <--
JP 3001919	B2	20000124		
PRAI JP 1990-3437		19900112		

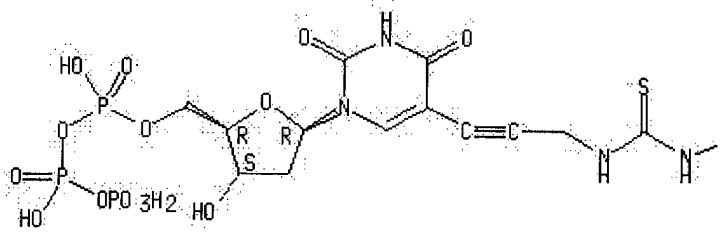
AB The **DNA** is amplified by PCR in the presence of ≥1 fluorescence-labeled nucleotides to prep. fluorescence-labeled **DNA** product, which is readily detectable without using radioactive substances. Amplification of human placenta **DNA** using c-Kr-ras/61 primers in the presence of fluorescence-labeled dUTP (FTC-AP-dUTP) was shown. The amplified **DNA** product was visibly detectable using immobilized oncogene probes (c-Ka-ras/61 Gly, Ser, and Val types).

IT **137993-59-0**
RL: PROC (Process)
(polymerase chain reaction in presence of, for easy visible detection of amplified product)

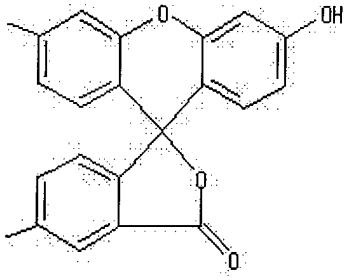
RN 137993-59-0 CAPLUS
CN Uridine 5'-(tetrahydrogen triphosphate), 2'-deoxy-5-[3-[[[(3',6'-dihydroxy-3-oxospiro[isobenzofuran-1(3H),9'-[9H]xanthen]-5-yl)amino]thioxomethyl]amino]-1-propynyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A



PAGE 1-B



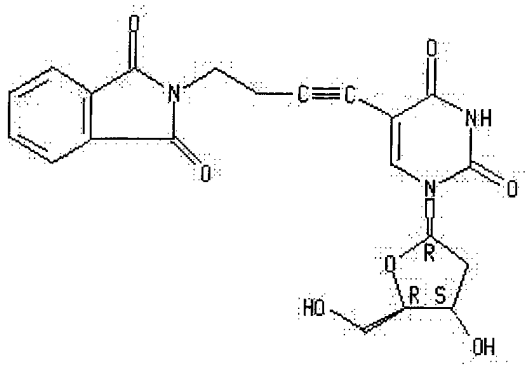
L5 ANSWER 6 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
Full Text

AN 1991:409259 CAPLUS
DN 115:9259
TI Preparation of crosslinking oligonucleotides as **nucleic** acid hybridization probes
IN Petrie, Charles R.; Meyer, Richard B.; Tabone, John C.; Hurst, Gerald D.
PA Microprobe Corp., USA
SO PCT Int. Appl., 42 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 8

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9014353	A1	19901129	WO 1990-US2740	19900515 <--
	W: CA, JP				
	RW: AT, BE, CH, DE, DK, ES, FR, GB, IT, LU, NL, SE				
	EP 472648	A1	19920304	EP 1990-908844	19900515
	R: AT, BE, CH, DE, DK, ES, FR, GB, IT, LI, LU, NL, SE				
	JP 04507402	T2	19921224	JP 1990-508242	19900515
	US 5824796	A	19981020	US 1994-334490	19941104
PRAI	US 1989-353857	A	19890518		
	US 1988-250474	B2	19880928		
	WO 1990-US2740	W	19900515		
	US 1993-49807	B1	19930420		

OS MARPAT 115:9259
AB R1-B-(CH2)q-Yr-(CH2)m-A1 [R1 = H, sugar (analog) moiety optionally contg. Q1, Q2, Q3, P, etc.; Q1 = OH, OP(O)(OH)2, OP(O)(OH)OP(O)(OH)2; Q2 = O, S; Q3 = CH2R2, SR2, OR2, NR2R3; R2, R3 = H, alkyl; B = **nucleic** acid base or an analog thereof; Y = functional linking group; m, q = 0, 1-8 integer; r = 0, 1; A1 = leaving group], useful as **nucleic** acid hybridization probes and therefore useful for diagnosis of diseases, were prepd. Reaction of 5-iodo-2'-deoxyuridine in DMF with 4-phthalimido-1-butyne in the presence of (Ph3P)4Pd and Et3N at 60° for 3 h gave 5-(4-phthalimido-1-butyn-1-yl)-2'-deoxyuridine, whose hydrogenation over Raney Ni gave 5-(4-phthalimidobutyl)-2'-deoxyuridine. 5-[3-(Trifluoroacetamido)propyl]-2'-deoxyuridine was prepd. similarly and converted according to known methods into 5'-O-(dimethoxytrityl)-2'-deoxyuridine-3'-(N,N-diisopropyl)phosphoramidite cyanoethyl ester, which was used in the automated synthesis of 3'-CT TCC U1TG TAG CTG-5' [I; U1 = 5-(3-aminopropyl)-2'-deoxyuridine residue]. This was reacted with N-(iodoactoxy)succinimide to give II [U1 = 5-(3-iodoacetamidopropyl)-2'-uridine residue], whose crosslinking to a 30-mer oligonucleotide derived from human papillomavirus (HPV) was evaluated.
IT **134140-85-5P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. and hydrogenation of)
RN 134140-85-5 CAPLUS
CN Uridine, 2'-deoxy-5-[4-(1,3-dihydro-1,3-dioxo-2H-isoindol-2-yl)-1-butynyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L5 ANSWER 20 OF 45 CAPLUS COPYRIGHT 2003 ACS on STN
Full Text
AN 1989:439836 CAPLUS
DN 111:39836
TI Preparation of 5-(3-aminoprop-1-ynyl)deoxyuridine derivatives and their use for synthesis of **DNA** and RNA labeled with nonradioactive markers.
IN Haralambidis, Jim
PA Florey, Howard, Institute of Experimental Physiology and Medicine, Australia
SO PCT Int. Appl., 58 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 8810264	A1	19881229	WO 1988-AU207	19880624 <--
	W: AU, JP, US				
	RW: AT, BE, CH, DE, FR, GB, IT, LU, NL, SE				
	AU 8819909	A1	19890119	AU 1988-19909	19880624 <--
	AU 598946	B2	19900705		
	EP 366685	A1	19900509	EP 1988-905594	19880624 <--
	EP 366685	B1	19941019		
	R: AT, BE, CH, DE, FR, GB, IT, LI, LU, NL, SE				
	JP 02504144	T2	19901129	JP 1988-505657	19880624 <--
	JP 2828642	B2	19981125		
	CA 1340032	A1	19980908	CA 1988-570427	19880624
PRAI	AU 1987-2666	A	19870624		
	WO 1988-AU207	A	19880624		

OS MARPAT 111:39836
AB The title nucleoside derivs. [I; Y = H, (un)protected OH; X = H, phosphonate group, P(OQ)NR1R2; R1, R2 = (un)branched, (un)substituted alkyl; Q = phosphate protecting group; Z = H, phosphate, triphosphate group; X1 = (un)branched (1-15 alkyl; R = amino protecting group, A, Y1NHA; A = fluorophore (e.g. fluorescein) or other non-radioactive detectable group (e.g. biotin, avidin, colloidal Au or Ag, ferritin, and enzymes such as β-galactosidase, urease, peroxidase); γ1 = (un)branched C1-10 alkylcarbonyl], useful for prepg. **DNA** and RNA labeled with non-radioactive detectable markers as **nucleic** acid hybridization probes, were prepd. Thus, coupling of 3',5'-di-O-p-toluoyl-5-iododeoxyuridine with BOCNHCH2C≡CH (BOC = CO2CMe3) in EtOAc in the presence of (Ph3P)2PdCl2, CuI and Et3N gave 84% I (Y = H, Z = X = toluoyl, X1NHR = C ≡ CCH2NHGBOC). Sapon. of the latter with K2CO3 in MeOH followed by reaction with 4,4'-dimethoxytrityl chloride (DMTrCl) in pyridine gave 68% I (X = Y = H, Z = DMTr, X1NHR = C≡CCH2NHBOC) which was treated with [(Me2CH)2N]2POMe in CH2Cl2 contg. tetrazole and (Me2CH)2NH to give I [Y = H, Z = CMTr, X = P(OMe)N(CHMe2)2, X1NHR =

C≡CCH2NHBOC]. The latter can be reacted by the phosphoramidite method to prep. oligonucleotides which incorporate the modified nucleoside on the 5'-end or internally.

IT **114079-33-3P**

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(prepn. and acylation of, with nitrophenylaminohexanoate deriv.)

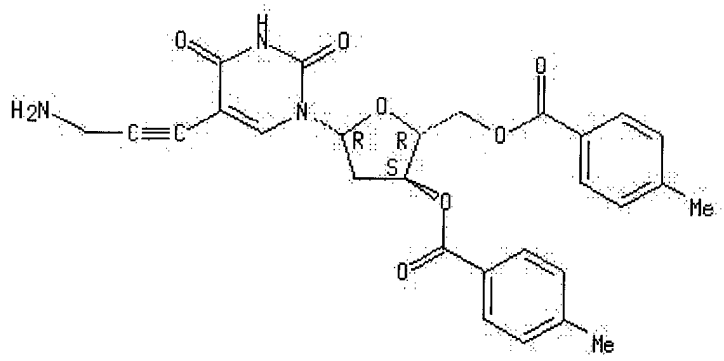
RN 114079-33-3 CAPLUS

CN Uridine, 5-(3-amino-1-propynyl)-2'-deoxy-, 3',5'-bis(4-methylbenzoate), mono(trifluoroacetate) (salt) (9CI) (CA INDEX NAME)

CM 1

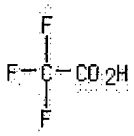
CRN 114079-32-2
CMF C28 H27 N3 O7

Absolute stereochemistry.



CM 2

CRN 76-05-1
CMF C2 H F3 O2



IT **114079-31-1P 114079-35-5P**

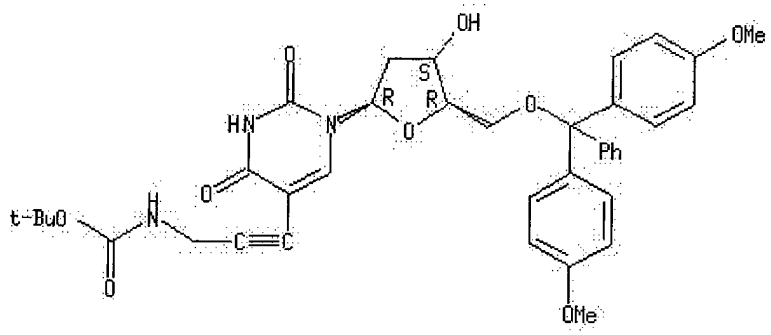
RL: SPN (Synthetic preparation); PREP (Preparation)

(prepn. and condensation of, with bis(diisopropylamino)methoxyphosphine)

RN 114079-31-1 CAPLUS

CN Carbamic acid, [3-[1-[5-O-[bis(4-methoxyphenyl)phenylmethyl]-2-deoxy-β-D-erythro-pentofuranosyl]-1,2,3,4-tetrahydro-2,4-dioxo-5-pyrimidinyl]-2-propynyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

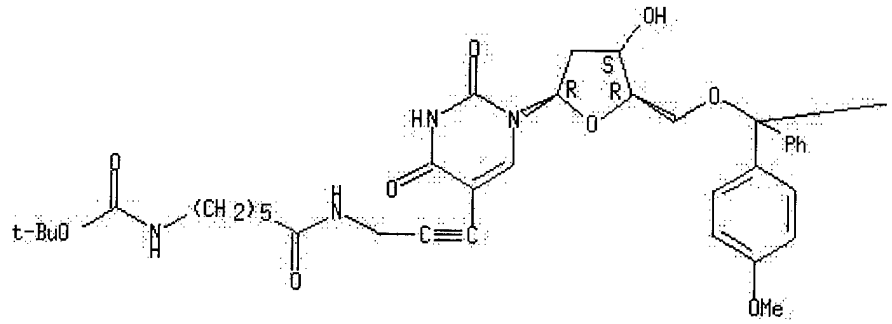
Absolute stereochemistry.



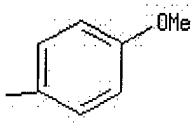
RN 114079-35-5 CAPLUS
CN Carbamic acid, [6-[[3-[1-[5-O-[bis(4-methoxyphenyl)phenylmethyl]-2-deoxy-β-D-erythro-pentofuranosyl]-1,2,3,4-tetrahydro-2,4-dioxo-5-pyrimidinyl]-2-propynyl]amino]-6-oxohexyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A

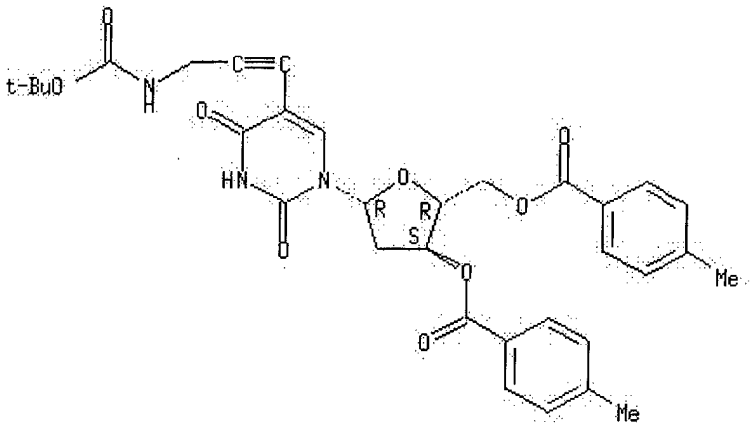


PAGE 1-B



IT **114079-29-7P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. and reaction of, in prepn. of **nucleic** acid marker)
RN 114079-29-7 CAPLUS
CN Carbamic acid, [3-[1-[2-deoxy-3,5-bis-O-(4-methylbenzoyl)-β-D-erythro-pentofuranosyl]-1,2,3,4-tetrahydro-2,4-dioxo-5-pyrimidinyl]-2-propynyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.



IT **114079-34-4P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. and sapon.-alkylation of, by dimethoxytrityl chloride)
RN 114079-34-4 CAPLUS
CN Carbamic acid, [6-[[3-[1-[2-deoxy-3,5-bis-O-(4-methylbenzoyl)-β-D-erythro-pentofuranosyl]-1,2,3,4-tetrahydro-2,4-dioxo-5-pyrimidinyl]-2-propynyl]amino]-6-oxohexyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

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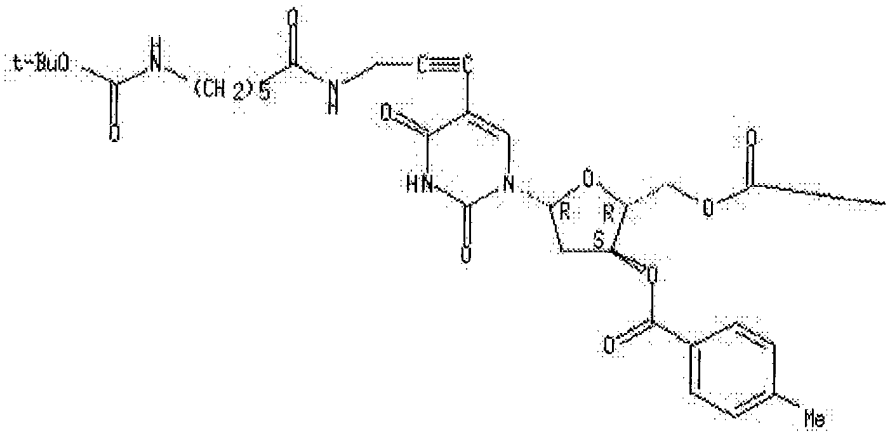

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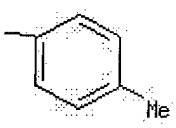

IT **114079-36-6P 114103-42-3P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. of, as intermediate for nonradioactive marker-labeled oligonucleotides)
RN 114079-36-6 CAPLUS
CN Carbamic acid, [6-[[3-[1-[5-O-[bis(4-methoxyphenyl)phenylmethyl]-3-O-[[bis(1-methylethyl)amino]methoxyphosphino]-2-deoxy-β-D-erythro-pentofuranosyl]-1,2,3,4-tetrahydro-2,4-dioxo-5-pyrimidinyl]-2-propynyl]amino]-6-oxohexyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

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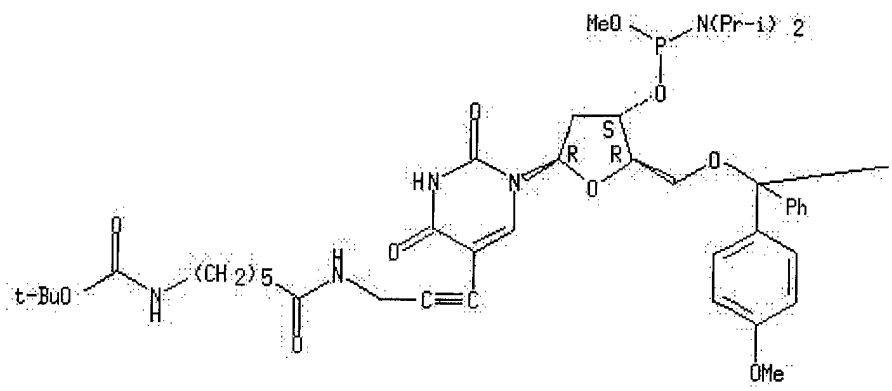

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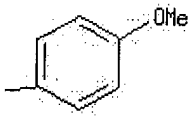
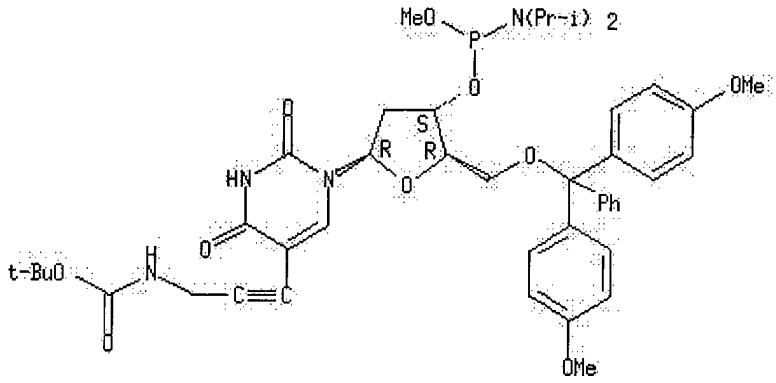


RN 114103-42-3 CAPLUS
CN Carbamic acid, [3-[1-[5-O-[bis(4-methoxyphenyl)phenylmethyl]-3-O-[[bis(1-methylethyl)amino]methoxyphosphino]-2-deoxy-β-D-erythro-pentofuranosyl]-1,2,3,4-tetrahydro-2,4-dioxo-5-pyrimidinyl]-2-propynyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

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FULL ESTIMATED COST                              48.59      197.35

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)    SINCE FILE      TOTAL
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CA SUBSCRIBER PRICE                              -3.91       -3.91

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